

# **HDA** Standard Pharmaceutical Product and Medical Device Information (Rx Product Only)

Version 2024					Introduction 7	ype: New Item	n		x Final Version			Date:	8/30/	2024			
PRODUCT INFORMATION								SPECIAL HANDLING AND STORAGE REQUIREMENTS*									
Company Name: Camber Pharmaceuticals, Inc.					Application: ANDA			a. Temperature – Indicate the USP temperature range for this product.									
Application Number for NDA/ANDA/BLA; PMA/510(k): 216591				NDA 505(b) Type: NOT APPLICABLE			Temperature Range Controlled Room – between 20 and 25 C (68° – 77° F)										
Medical Device Class, if applicab	le:																
	11-856-3719									-	Other Temperature Range I	Requirement	After reconstitution	, this vial may be s	ored in a refrigerate	or for 96 hours	
Proprietary Name (If Applicable) ar		ime:		drochloride for Inject	ion, USP 500 i	mg/vial (Single					(write in)		without significant	loss or potency.			
	31722-210-10			Unit of Use NDC:			UPC:	331722210102			Notes						
UDI				CVX Code:			MVX Code:										
Description: Vancomycin Hydrochloride for Injection, USP 500 mg/vial (Single-dose Vial)									Is this product to be shipped				No				
										Is this product to be shipped	d to customers on d	ry ice?		No			
Active Ingredient(s):  Vancomycin hydrochloride, USP										b. Contact for temperature excursion questions:							
URL for Additional Product Information: www.camberpharma.com										Name: Soma Raju							
Address:	800 Centennial Ave, Suite 1				Address 2:				Number: 732-529-0423								
City:	Piscataway					State:	- Lip. 0000 i			Group E-mail: somaraju@heterousa.com							
Key Contact:	Customer Service					Email:					•						
Phone Number:	1-866-827-3647	47				Fax: 732-562-8788				c. Special regulations for product in any states?					No		
Product Therapeutic Classification	roduct Therapeutic Classification: Tricyclic glycopeptide antibiotic									Special returns requirements for this product?							
ADDITIONAL PRODUCT INFORMATION PRODUCT DESCRIPTION INFORMATION d. Store product (unit of sale) upright?																	
	ADDITI	ONAL PRODU				PRODUCT DESCRIPTION INFORMATION					ct (unit of sale) upright?		No				
The product is?				Product	Direct-Ship C	nly	1				Protect product (unit of sa	ale) from light?			No		
a legend device?		No		Product	Unit Dose		Size:	10 single dose via	als	e. Shelf life:					24	Months	
if yes, enter class #		NI.	Orph	an Drug Status				500 / -		1	Initial shelf life at launch (	if different):				Months	
a product kit? if yes, list NDCs of		No FDA Approval Status				Strength:	500 mg/vial				ORDER INFORM	IATION					
component parts			FDA	Approvai Status				Sterile, lyophilized powder	r or cake for			ORDER IIII ORII	ATION				
reverse numbered?		No					Dosage Forn	<ol> <li>preparing intravenous infu injections</li> </ol>	usions or		Unit of Sale		What is the	NDC selling	unit?		
co-licensed?		No	Allerg	gens Present				mecauris			Bottle		1 Box of 10	/ials			
latex-free?		Yes					Product Sha	Single dose vial			x Box/Carton		(Write-in, e.	g. 1 Box of 10	Vials)		
preservative-free?		Yes					1 Todact Ond				Ampule						
correctional institution block?		No					Product Cole	White to tan			x Glass		Minimum or	der quantity	?	Yes	
opioid?		No		to a f O dada	India			NI/A			Tube						
Cannabinoid?  If Unit Dose, is item bar coded to un	nit does for	No	Coun	try of Origin	inuia		Product Imp	rint: N/A			Vial Liquid Sgl Vial Liquid Multi		If Voc. how	many of whi	ch nackado t	hyno?	
hospital scanning?	THE GOSE TO	Yes	Is this	s product covered un	der the					Vial Liquid Multi If Yes, how many of which package type?  x Vial Powder Sgl 1 Each					ype:		
If Unit Dose, indicate NDC here:		31722-210-3		Agreements Act (TA		No				Vial Powder Multi Inner/Carton/Pack							
11 Elik Bese, indicate 1150 fictor								Other: Write In Case									
			FOR G	ENERIC DRUG PRO	DUCTS												
					-	Au	thorized Generic	*If Authorized Generic, or				IARMACY ORDER					
I. Orange Book Rating:					section fields are not applicable			Rec. sell unit to customer? Rx billing unit to pharmacy:									
II. Generic Equivalent to What Brand?: Vancomycin Hydrochloride for Injection (Fresenius Kabi USA, LI				LC)				Each									
DRUG SUPPLY CHAIN SECURITY ACT (DSCSA) INFOR				RMATION				(Write-in, e.g. 1 Vial)  HCPCS J-Code:  Gram  Milliliter									
										111111111111111111111111111111111111111	J3370	1					
Does supplier meet DSCSA definit	ion of manufactur	er?		Yes		GLN:	0331722498975					AND PACKING IN	NFORMATION	I			
Is product exempt from DSCSA?			No														
If yes, select exemption:						GCP:					Weight Lbs.	Dimensi	ons (US msm	ıts.)		Saleable #	
Other exemption - Write in:											Weight Lbs.	Depth	Width	Height	(Cube)	Pieces	
Is product repackaged?			No	Voc			iginal product pur	chased		Item/Each:	0.5	6.31	2.5	2.94	46.38	1	
Is product sold by manufacturer's Has FDA granted waiver/exception				Yes No	-	direct from m		r repackaged product		Box/Carton/Bu	n-41a/						
If yes, attach documentation from		oduct?		110	1	Provide sour	ce manuracturer ro	г гераскадео ргооист		Inner Pack:	naie/						
, , , , , , , , , , , , , , , , , ,										Case:	44.05	13,25	13.38				
			GTIN AND H	IIBCC PRODUCT IN	FORMATION						11.65	13.25	13.38	6.63	1175.40	20	
										Pallet:							
Saleable Unit of Measure	RFID tag(Y/N)	Saleable	HIBC	C		GTI	N-14	Unit of Use GTIN	N-14								
		Quantity				002	24722240402										
x       ltem/Each   N   1					00331722210102			COST INFORMATION WHOLESALER USE ONLY:					Y:				
X Case	N	20				203	31722210106	-									
Pallet								1		Regular Cost			Vendor #:				
										Invoice Cost (V	VAC) (\$)	\$30.00	Whsl. Code				
													Fineline Co	de:			
										As of date:	1/30/2023		ļ				
<u> </u>			Attach o	CODY of SAFETY DAT	A SHEET (SD	S) or non haza	rd letter PACKACE	INSERT, LABEL AND PH	IOTO OE PE	RUDITICT BYCKYO	SING and RAPCODE		I				
					A SHILL I (SD	U) UI HUH HAZA	IG IGHEI, I AGNAGE	INOLINI, LADEL AND FR	IO O OI FI	CODOCT ACKAC	JING AND DANGODE.						



## **Standard Pharmaceutical Product and Medical Device Information (Rx Product Only)**

#### Version 2024

For Designated Drop Ship Only Products, Please Use Page 3

MATERIAL HAZ	ARD CLASSIFICATION and TRANSPORTATION							
Is this product (check all that apply):								
a. Cytotoxic?	SDS Hazard Classification							
b. CA Prop. 65 Carcinogen or Reproductive Toxicant?	1							
Is the product a CA Prop 65 carcinogen?	x Organic Corrosive							
Is the product a CA Prop 65 reproductive toxicant?	Inorganic Oxidizer							
Does the product label bear a CA Prop 65 warning?	Steroid/Androgen Contact Hazard							
c. Contact Hazard?	Does the product have an Aerosol class? If yes, No							
d. Does this product require special clean-up instructions?	identify NFPA Storage Level:							
(If yes, attach SDS with special instructions.)	NFPA Storage Level:							
e. Does the product contain DEHP?								
Is this product regulated for shipment by DOT?	Is the product a NIOSH hazardous drug?							
(if yes, answer a-e below and provide SDS)	If yes, indicate which:							
a. UN/Identification Number	ii yes, indicate which.							
b. Proper Shipping Name								
c. DOT Hazard Class	Hazardous Waste Identification							
d. Packing Group								
e. Inhalation Hazard?	EPA Hazardous Waste Code: Waste Characteristics							
, ,	DEMS of DECISTOR DESTRICTIONS							
(if yes, answer a-e below and provide SDS) a. UN/Identification Number	REMS or REGISTRY RESTRICTIONS							
	Is there a REMS on this product?							
b. Proper Shipping Name c. DOT Hazard Class	If Yes, is it managed with a pharmacy registry?							
d. Packing Group	Website URL:							
e. Inhalation Hazard?	Website UKL.							
Is the product restricted for air shipment? If so, indicate restriction:	Med Guide Required No							
Passenger	Limited Distribution Requirement							
Cargo	Comments / Details: (For example, iPledge program?)							
Passenger & Cargo								
Is this a reportable quantity? No	REMS: No							
RQ Threshold:	REMS Program Manager Name: Phone:							
Is this a marine pollutant? No	Supplier Manages REMS registry exclusively:							
Is this product shipped utilizing an authorized DOT exception or Special Permit?	Wholesale distributor support:							
No (if yes, identify method below)	Provider Name: DEA #:							
Limited Quantity	Site Enrollment Number assigned NCPDP#:							
Consumer Commodity, ORM-D	by Supplier: NPI #:							
Small Quantity (49 CFR 173.4)	Comments							
Special Permit; DOT-SP	Comments							
Special Provision (listed in Column 7 of 49 CFR 172.101);								
SP#	Registry: No							
ADDII CTODAGE INFORMATION	Registry Program Contact Name: Phone:							
ADD'L STORAGE INFORMATION	Comments							
Is the Product								
Controlled Substance? No Controlled Substance Code	RETURN INSTRUCTIONS							
Controlled by State(s)? No Listed Chemical (List I or II) No								
ARCOS Reportable? No If yes, indicate which:	Contact tel. # if product received damaged: 1-866-827-3647							
Schedule No. Is it a scheduled listed chemical product?: No	Is product returnable for credit:							
CLASS OF TRADE RESTRICTION:	URL/Link to returns policy:							
No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and physician offices  Yes	contact - customerservice@camberpharma.com							
Restricted to retail pharmacy only:  No								
	Special regulations or returns requirements for this product in certain states?							
Restricted to hospital, clinics, and physician offices only:  No	, INO							
Restricted from US territories? (explain in comments)	If so, which states? Other requirements? Comments?							
Comments:								
MISCELLANEO	DUS NOTES and/or Image of Product Barcode:							



## **Standard Pharmaceutical Product and Medical Device Information (Rx Product Only)**

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### FOR DESIGNATED DROP SHIP PRODUCT ONLY - if not a designated drop ship, do not complete.

Order Method for Designated Drop Ship Product	Standard Order Receipt and Processing							
Purchase orders may be accepted by:  a. EDI  b. Autofax  c. Fax  d. Phone only e. Supplier Web Site only  Minimum Order Quantity:  Supplier's Customer Service Number:  Contracted 3PL company / contact #:  Name: Phone:	Purchase order daily receipt cut off time by supplier Cut off time:  Shipping lead time of PO: Hours Days Ships same day for next day receipt: Ships for second day receipt: Ships regular ground for 3-10 days receipt:							
Expedited Freight Charges or Other Designated Drop Ship Fees:	Overnight and Priority Overnight PO Processing							
Expedited freight fees billed with each order:  Drop Ship service fee billed with each order:  Drop Ship miscellaneous fees billed:  Comments:	Overnight receipt available:  PO Receipt cut off time:  Days of week overnight is available:  Monday Tuesday Wednesday Thursday Friday							
	Priority Overnight receipt available:							
Class of Trade Restriction:  No restriction: Select YES if sold to retail pharmacy, hospitals, clinics and physician offices Restricted to retail pharmacy only: Restricted to hospital, clinics, and physician offices only: Restricted from US territories? (explain in comments)  Comments:	PO Receipt Cut off time:  Saturday Overnight receipt available: PO Receipt Cut off time: Phone: Phone: Fax: EDI: Overnight Fees apply: Other fees apply:							
Other Data Information Required to Process PO:	Return Instructions							
Patient Procedure Date: Physician Name: Physician/Clinic Phone # Physician State License # Physician/Clinic DEA #: Physician/Clinic Specialty:  Miscellaneous Notes:	Contact # if product is received damaged: Is product returnable for credit: URL/Link to returns policy:  Special regulations or returns requirements for this product in certain states? If so, which states? Other requirements? Comments?							
	ADDITIONAL INFORMATION							
	Is product order for scheduled patient procedure? Is product order for restocking purposes?							